

Utah Medicaid Provider Manual	Drug Criteria and Limits
Division of Health Care Financing	Updated January 2009

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Explanation of Medicaid Policy

Drugs with Criteria and Limits

Many drugs in the Medicaid pharmacy program do not require a Prior Authorization (PA), but are still subject to restrictions that are outlined in the Medicaid Pharmacy Services Manual and the Medicaid Physician Services Manual. This section serves as a quick reference for the specific policies that govern coverage of these drugs.

In accordance with the Utah Medicaid Provider Manual for Pharmacy Services, SECTION 2, Chapter 4-9, Limits on Certain drugs, some drugs are limited by a quantity in any thirty-day period. These drugs do not qualify for early refills, as stated in Chapter 4-7, Early Refills. The limits listed are those approved by the Medicaid Drug Utilization Review (DUR) Board. Physicians and other prescribers who feel that a patient has specific needs that exceed the limits may appeal to the DUR Board. All medications remain subject to all the other requirements of the Utah Medicaid Pharmacy Program, as described in the Utah Medicaid Manual for Pharmacy Services.

Drugs Requiring Prior Authorization

In accordance with the Utah Medicaid Provider Manual for Pharmacy Services, SECTION 2, Chapter 3, certain drugs that are covered by the Medicaid program may require the patient and physician to meet specific criteria and demonstrate medical necessity in order to receive the requested medication. Detailed information regarding prior approval criteria for individual medications and classes of medications is provided in this manual.

Please note that prior authorization for a medication is client specific, pharmacy specific, and product specific. Prior authorization cannot be transferred to another pharmacy, to another product, nor to another strength of a product that has been approved. The prior authorization cannot be transferred to another client.

To initiate a prior authorization request, the physician must gather all of the records that are requested in the criteria set for the medication being prescribed. These records should then be faxed, along with a cover sheet that includes the client's name and client ID, physician's name and telephone number, and (if known) the name and telephone number of the pharmacy that the client would like to use. A fax cover sheet that can be filled out with the requested information is included in the back of the prior authorization section, should you wish to use it. The requests can be faxed to (801) 536-0477.

All injectable products, with the exception of 10ml vials of insulin, require prior authorization under the Non-Traditional Medicaid plan.

Non-Traditional Medicaid has additional restrictions in place. In accordance with the Utah Medicaid Provider Manual for Non-Traditional Medicaid, SECTION 2, Chapter 2-19.2, no lozenges, suckers, rapid dissolve, lollipop, pellets, patches or other unique formulation delivery methodologies developed to garner "uniqueness" will be covered, except where the specific medication is unavailable in any other form (Duragesic and Actiq). Drugs are covered for labeled indications only.

Exceptions to Policy

All requests for exceptions to policy require a petition to the DUR board. DUR meetings are held on the second Thursday of every month. Petitions to the DUR board must be received one week prior to the monthly meeting. Petitions may be faxed to the prior authorization team.

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Drugs with Criteria and Limits	
<p>ADD/ADHD Medications</p> <ul style="list-style-type: none"> Amphetamines Methylphenidate & Derivatives Strattera 	<p>Amphetamines:</p> <ul style="list-style-type: none"> Age 0-2: Not a covered benefit. Age 3-5: Immediate-release Adderall and Dexedrine generic formulations are covered - Valid ICD-9 code must be written on the prescription. Age 6-18: Covered - Valid ICD-9 code must be written on the prescription. Age 19+: Prior Authorization Required - see page 21. <p>Methylphenidate & Derivatives:</p> <ul style="list-style-type: none"> Age 0-5: Not a covered benefit Age 6-18: Covered - Valid ICD-9 code must be written on the prescription. Age 19+: Prior Authorization Required - see page 21. Daytrana patch is non-covered under Non-Traditional Medicaid <p>Strattera:</p> <ul style="list-style-type: none"> Covered for ages 6+. Cumulative limit of 66 capsules in 30 days. Approved as a stand-alone treatment for ADHD.
<p>Analgesics</p> <ul style="list-style-type: none"> Celebrex Tramadol/Ultracet Fentanyl Patch Fentora Actiq Methadone Long-Acting Opioids (Avinza, Kadian, MS-Contin, Oxycontin & generics) Short-Acting Opioids Short-Acting Opioid/APAP 	<p>Celebrex:</p> <ul style="list-style-type: none"> Age below 65: Prior Authorization Required - see page 12. Age 65+: Cumulative limit of 60 capsules in 30 days. <p>Tramadol/Ultracet:</p> <ul style="list-style-type: none"> Cumulative limit of 180 tablets in 30 days <p>Fentanyl Patch</p> <ul style="list-style-type: none"> Cumulative limit of 15 patches in 30 days The cumulative limit may be overridden if the prescriber provides a valid ICD-9 diagnosis code for cancer. The 100mcg Fentanyl Patch is <i>only</i> covered with valid ICD-9 diagnosis code for cancer. <p>Fentora</p> <ul style="list-style-type: none"> Cumulative limit of 120 units in 30 days - Actiq counts towards this limit. Only a covered benefit if the prescriber provides a valid ICD-9 diagnosis code for cancer. <p>Actiq</p> <ul style="list-style-type: none"> Cumulative limit of 120 units in 30 days - Fentora counts toward this limit. Only a covered benefit if the prescriber provides a valid ICD-9 diagnosis code for cancer. <p>Methadone</p> <ul style="list-style-type: none"> Cumulative limit of 150 tablets in 30 days. The cumulative limit may be overridden if the prescriber provides a valid ICD-9 diagnosis code for cancer. <p>Long-Acting Opioids</p> <ul style="list-style-type: none"> Cumulative limit of 90 tablets in 30 days. The cumulative limit may be overridden if the prescriber provides a valid ICD-9 diagnosis code for cancer. <p>Short-Acting Opioids</p> <ul style="list-style-type: none"> Cumulative limit of 180 tablets in 30 days The cumulative limit may be overridden if the prescriber provides a valid ICD-9 diagnosis code for cancer. <p>Short-Acting Opioid/APAP Combinations</p> <ul style="list-style-type: none"> Cumulative limit of 180 tablets in 30 days The control is on Acetaminophen and may not be overridden for safety reasons.

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Drugs with Criteria and Limits	
Atypical Antipsychotics (Abilify, Clozaril, Geodon, Invega , Risperdal, Seroquel, Sybmbyax, Zyprexa)	<ul style="list-style-type: none"> Valid ICD-9 diagnosis code is required on each prescription. ICD-9 codes may be found in the <u>Utah Medicaid Provider Manual for Physicians Services and Anesthesiology</u>. ICD-9 code must be correct for the patients age. Risperdal Consta requires a prior authorization - see page 21.
Benzodiazepines	<ul style="list-style-type: none"> Cumulative limit of 120 tablets/capsules in 30 days. Short acting benzodiazepines that are typically used to treat insomnia are governed by the criteria for sedative-hypnotics. Prior Authorization required for Xanax XR - see page 22. Xanax XR is non-covered under Non-Traditional Medicaid.
Bupropion (Zyban, Wellbutrin)	<ul style="list-style-type: none"> One of two valid ICD-9 diagnosis codes is required on each prescription. ICD-9 311 indicates depressive disorders. ICD-9 305.1 indicates smoking cessation. Wellbutrin XL is non-covered under Non-Traditional Medicaid.
Butalbital Containing Products	<ul style="list-style-type: none"> Cumulative limit of 30 tablets in 30 days.
*	
Cymbalta	<ul style="list-style-type: none"> One of two valid ICD-9 diagnosis codes is required on each prescription. ICD-9 311 indicates depressive disorders. ICD-9 729.2 for neuralgias, etc. The maximum daily dose is 60 mg. Monthly quantity limits are set accordingly.
Diphenoxylate Containing Products	<ul style="list-style-type: none"> Cumulative limit of 30 tablets in 30 days.

Inhalers	LIMIT IN ANY 30 DAY PERIOD				
Effective April 1, 2002, the cumulative number of inhalers in any 30-day period is limited for a Medicaid client. The limit is set by class (excepting Foradil and Serevent which are limited by NDC number). This means the highest number in any one class is the maximum. When there are more than two sizes or strengths for a given product, the limit is based on the largest size or strength. There are two groups of inhalers: oral and nasal. For each group, the limits are stated below.					
Inhaler Class	Generic Name	Brand Name	Product Size	Doses per Inhaler	Maximum No. In 30 Days
Nasal Anti-inflammatory inhalers	beclomethasone	Beconase AQ	25	200	2
	fluticasone	Flonase	16	120	1
	triamcinolone	Nasacort AQ	16.5	120	2
	triamcinolone	Nasacort HFA	9.3	100	3
	flunisolide	Nasarel	25	200	3
	mometasone	Nasonex	17	120	1
	budesonide	Rhinocort AQUA	8.4	120	2

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Inhaler Class	Generic Name	Brand Name	Product Size	Doses per Inhaler	Maximum No. In 30 Days
Beta 2 agonists and Sympathomimetic Inhalers	Albuterol	generic	17 gm	200	4
		Proventil	17 gm	200	4
		Proventil HFA	6.7 gm	200	4
		Ventolin	6.8 gm	80	4
			17 gm	200	4
		Ventolin HFA	18gm	200	4
	Formoterol	Foradil		12	1
				60	2
	Metaproterenol	Alupent	14 gm	200	2
	Pirbuterol	Maxair	25.6 gm	300	3
	Pirbuterol	Maxair Autohaler	14 gm	400	1
	Salmeterol	Serevent	6.5 gm	60	1
			13 gm	120	1
			Serevent Diskus	60	1
Anticholinergic Inhalers	Ipratropium	Atrovent HFA	14 gm	200	2
	Ipratropium / Albuterol	Combivent	14.7 gm	200	2
	Tiotropium	Spiriva	30 cap.	30	1
Anti-Inflammatory Inhalers	Beclomethasone	Qvar 40mg	7.3 gm	100	2
		Qvar 80mg	7.3gm	100	2
	Budesonide	Pulmicort Turbuhaler		200	2
	Flunisolide	AeroBid, AeroBid-M	7 gm	100	2
			13 gm	120	1
				120	1
				120	2
	Fluticasone DPI	Flovent Rotadisk 50 mcg, 100 mcg, and 250 mcg		60	1
				60	1
				60	4
	Triamcinolone MDI	Azmecort	20 gm	240	2
				60	1
				60	1
				60	1
	Fluticasone / Salmeterol DPI	Advair diskus 100/50		60	1
				60	1
				60	1
Mast cell stabilizer Inhalers	Cromolyn MDI	Intal	8.1 gm	112	3
			14.2 gm	200	2
	Nedocromil MDI	Tilade	16.2 gm	112	3

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Drugs with Criteria and Limits													
Laxatives <ul style="list-style-type: none"> Miralax Lactulose 	Miralax: <ul style="list-style-type: none"> Cumulative limit of 1054gm in 30 days Miralax OTC formulations are not covered. Lactulose: <ul style="list-style-type: none"> Cumulative limit of 6,000ml in 30 days Over 6,000ml in 30 days requires a prior authorization - see page 16. 												
Levothyroxine Products	<ul style="list-style-type: none"> Generic use mandated when AB-rated equivalent exists Use the table below to determine appropriate substitutions: <table> <tr> <th><u>Drug</u></th><th><u>Rating</u></th></tr> <tr> <td>Unithroid</td><td>AB1,AB2, AB3</td></tr> <tr> <td>Mylan Levothyroxine</td><td>AB1,AB2,AB3</td></tr> <tr> <td>Levoxyl</td><td>AB1, AB3</td></tr> <tr> <td>Synthroid</td><td>AB2</td></tr> <tr> <td>Levo-T</td><td>AB2, AB3</td></tr> </table>	<u>Drug</u>	<u>Rating</u>	Unithroid	AB1,AB2, AB3	Mylan Levothyroxine	AB1,AB2,AB3	Levoxyl	AB1, AB3	Synthroid	AB2	Levo-T	AB2, AB3
<u>Drug</u>	<u>Rating</u>												
Unithroid	AB1,AB2, AB3												
Mylan Levothyroxine	AB1,AB2,AB3												
Levoxyl	AB1, AB3												
Synthroid	AB2												
Levo-T	AB2, AB3												
Migraine Medications (Triptans) <i>(Imitrex, Zomig, Amerge, Axert, Maxalt)</i>	<ul style="list-style-type: none"> Cumulative limit of 9 dosage units per 30 days - all forms count towards this limit. Examples of drugs in this class include Imitrex, Maxalt, and Zomig. 												
Muscle Relaxants	<ul style="list-style-type: none"> Cumulative limit of 30 tablets in 30 days. Dantrolene, Baclofen, and Tizanidine are not included in this policy. 												
Prograf (tacrolimus)	<ul style="list-style-type: none"> All <u>oral</u> dosage forms are a covered benefit for use as a prophylaxis of organ rejection in allogenic liver transplants only. All injectable dosage forms are covered in physician office or hospital only. 												
Proton Pump Inhibitors	<ul style="list-style-type: none"> Cumulative limit of 30 units in 30 days. Prior Authorization required for twice daily dosing - see page 18. Prilosec OTC prescriptions do not require a PA for twice daily dosing. 												
Sedative-hypnotics for sleep <i>(Dalmane, Sonata, Somnote, Halcion, Ambien, Doral, Restoril, Lunesta, Rozerem, and their generics)</i>	<ul style="list-style-type: none"> Cumulative limit of 30 units in 30 days. Benzodiazepines that are typically used to treat insomnia are considered part of this class. 												

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- Minimum age requirement: 18 years old.
- Documented diagnosis of moderate to severe chronic plaque psoriasis.
- History of incomplete response or intolerance to at least one appropriate systemic agent or photo therapy.
- Negative TB skin test or history of treatment for latent TB infection.
- Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition.
- Dermatology consultation within the last 60 days.
- To be given in clinic setting only. Provider will bill with code J3490 and PA number.
- Patients with HMO's (except Select Health) will have to make arrangements with their HMO for coverage.
- Initial authorization is for 12 weekly injections.
- Additional 12 week course may be initiated provided CD4+T lymphocyte counts are within normal range and a minimum of 12 weeks have passed since the previous course of treatment.
- Maximum annual coverage is 24 weeks.

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<ul style="list-style-type: none"> Arava Enbrel 	<p>Arava:</p> <ul style="list-style-type: none"> Documented severe rheumatoid arthritis. Documented history of treatment, incomplete response, or intolerance to Methotrexate. Documented 6 or more swollen joints and 9 or more tender joints. Documented rheumatology consultation within the last 60 days. May not be given with other biological agents such as interferon, experimental medications or combinations. Initial authorization is for 6 months. Subsequent PA is for 12 months if the patient has at least 20% documented improvements in 4 of the following 6 areas: tender and swollen joint count, patient and/or global assessment of disease activity, pain, acute phase reactants. <p>Enbrel for Rheumatoid Arthritis or Plaque Psoriasis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Diagnosis of moderate to severe rheumatoid arthritis or plaque psoriasis. History of treatment, incomplete response, or intolerance to methotrexate or one other DMARD or second line drug (i.e. azathioprine, sulphadiazine, leflunomide, penicillamine, hydroxychloroquine, etc.) The number of swollen joints must be 6 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). The number of tender joints must be 9 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Enbrel may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Enbrel for Ankylosing Spondylitis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of ankylosing spondylitis. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Enbrel may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Enbrel for Juvenile Idiopathic Arthritis:</p> <ul style="list-style-type: none"> Minimum age requirement: 2 years old. Diagnosis of juvenile idiopathic arthritis. Documentation of failed treatment on at least one DMARD. Enbrel may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Enbrel for Plaque Psoriasis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of moderate to severe chronic plaque psoriasis. History of incomplete response or intolerance to at least one appropriate systemic agent or photo therapy. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Dermatology consultation within the last 60 days. Enbrel may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.

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Drugs Requiring Prior Authorization	
<ul style="list-style-type: none"> Humira 	<p>Humira for Juvenile Idiopathic Arthritis:</p> <ul style="list-style-type: none"> Minimum age requirement: 4 years old. Diagnosis of juvenile idiopathic arthritis. Documentation of failed treatment on at least one DMARD. Humira may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Humira for Crohn's Disease:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Diagnosis of moderate to severely active Crohn's Disease. Documented inadequate response to conventional therapy (i.e. 5-aminosalicylates, antibiotic, MTX, 6-mercaptopurine, azathioprine, corticosteroids, or budesonide). OR Documented intolerance to or loss of response on infliximab (Remicade). Humira may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Initial authorization is given for 1 year, and includes one 6-syringe starter pack and 2-syringe maintenance packs monthly thereafter. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Humira for Rheumatoid Arthritis or Plaque Psoriasis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Diagnosis of moderate to severe rheumatoid arthritis or plaque psoriasis. History of treatment, incomplete response, or intolerance to methotrexate or one other DMARD or second line drug (i.e. azathioprine, sulphadiazine, leflunomide, penicillamine, hydroxychloroquine, etc.) The number of swollen joints must be 6 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). The number of tender joints must be 9 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Humira may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Humira for Ankylosing Spondylitis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of ankylosing spondylitis. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Humira may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Humira for Plaque Psoriasis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of moderate to severe chronic plaque psoriasis. History of incomplete response or intolerance to at least one appropriate systemic agent or photo therapy. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Dermatology consultation within the last 60 days. Enbrel may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.

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<ul style="list-style-type: none"> Kineret 	<p>Kineret:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Diagnosis of moderate to severe rheumatoid arthritis. History of treatment, incomplete response, or intolerance to methotrexate or one other DMARD or second line drug (i.e. azathioprine, sulphadiazine, leflunomide, penicillamine, hydroxychloroquine, etc.) The number of swollen joints must be 6 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). The number of tender joints must be 9 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Rheumatology consultation within the last 60 days. Kineret may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.
<ul style="list-style-type: none"> Raptiva 	<p>Raptiva:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of moderate to severe chronic plaque psoriasis. History of incomplete response or intolerance to at least one appropriate systemic agent or photo therapy. Negative TB skin test or history of treatment for latent TB infection. Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. Dermatology consultation within the last 60 days. Raptiva may not be given with other biologic agents such as Interferon, experimental medications, or combinations. Initial authorization is given for 1 year. Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.

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<ul style="list-style-type: none"> • Remicaid 	<p>Remicaid for Crohn's Disease:</p> <ul style="list-style-type: none"> • Minimum age requirement: 6 years old. • Diagnosis of moderate to severely active Crohn's Disease. • Documented inadequate response to conventional therapy (i.e. 5-aminosalicylates, antibiotic, MTX, 6-mercaptopurine, azathioprine, corticosteroids, or budesonide). • Remicaid may not be given with other biologic agents such as Interferon, experimental medications, or combinations. • Remicaid may not be given with Enbrel or Kineret. • Negative TB skin test or history of treatment for latent TB infection. • Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. • To be given in clinic setting only and billed using code J1745 and PA number. • Patients in HMO's (other than Select Health) will need to make arrangements with their HMO for coverage. • Initial authorization is for 1 year. • Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Remicaid for Plaque Psoriasis:</p> <ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Documented diagnosis of moderate to severe chronic plaque psoriasis. • History of incomplete response or intolerance to at least one appropriate systemic agent or photo therapy. • Negative TB skin test or history of treatment for latent TB infection. • Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. • Dermatology consultation within the last 60 days. • Remicaid may not be given with other biologic agents such as Interferon, experimental medications, or combinations. • Remicaid may not be given with Enbrel or Kineret. • To be given in clinic setting only and billed using code J1745 and PA number. • Patients in HMO's (other than Select Health) will need to make arrangements with their HMO for coverage. • Initial authorization is for 1 year. • Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Remicaid for Rheumatoid Arthritis or Psoriatic Arthritis:</p> <ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Diagnosis of moderate to severe rheumatoid arthritis or plaque psoriasis. • History of treatment, incomplete response, or intolerance to methotrexate or one other DMARD or second line drug (i.e. azathioprine, sulphadiazine, leflunomide, penicillamine, hydroxychloroquine, etc.) • The number of swollen joints must be 6 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). • The number of tender joints must be 9 or more (WRITE SPECIFIC NUMBER IN NOTES OR LETTER). • Negative TB skin test or history of treatment for latent TB infection. • Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. • Rheumatology consultation within the last 60 days. • Remicaid may not be given with other biologic agents such as Interferon, experimental medications, or combinations. • Remicaid may not be given with Enbrel or Kineret. • To be given in clinic setting only and billed using code J1745 and PA number. • Patients in HMO's (other than Select Health) will need to make arrangements with their HMO for coverage. • Initial authorization is for 1 year. • Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.

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	<p>Remicaid for Ankylosing Spondylitis:</p> <ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Documented diagnosis of ankylosing spondylitis. • Negative TB skin test or history of treatment for latent TB infection. • Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. • Rheumatology consultation within the last 60 days. • Remicaid may not be given with other biologic agents such as Interferon, experimental medications, or combinations. • Remicaid may not be given with Enbrel or Kineret. • To be given in clinic setting only and billed using code J1745 and PA number. • Patients in HMO's (other than Select Health) will need to make arrangements with their HMO for coverage. • Initial authorization is for 1 year. • Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication. <p>Remicaid for Ulcerative Colitis:</p> <ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Diagnosis of moderate to severe ulcerative colitis. • Documented inadequate response to conventional therapy (i.e. 5-aminosalicylates, antibiotic, MTX, 6-mercaptopurine, azathioprine, corticosteroids, or budesonide). • Remicaid may not be given with other biologic agents such as Interferon, experimental medications, or combinations. • Remicaid may not be given with Enbrel or Kineret. • Negative TB skin test or history of treatment for latent TB infection. • Absence of active bacterial or viral infection, malignancy, or immunosuppressive condition. • To be given in clinic setting only and billed using code J1745 and PA number. • Patients in HMO's (other than Select Health) will need to make arrangements with their HMO's for coverage. • Initial authorization is for 1 year. • Re-authorization requires an updated letter of medical necessity or progress notes showing improvement or maintenance on medication.
Avastin	<ul style="list-style-type: none"> • Minimum age - 18 years old. • Documentation of diagnosis of metastatic carcinoma of colon or rectum OR non-squamous, non-small cell lung cancer OR macular degeneration. • Initial authorization may be granted for 1 year - renewal requires an updated letter of medical necessity.
Betamethasone Topical (Luxiq, Olux)	<ul style="list-style-type: none"> • Documented failure on generic formulations of betamethasone valerate creams or ointments within the last 12 months. • Initial authorization is given for 6 months. Subsequent authorizations require a telephone request from the physician's office or pharmacy.
Botox	<ul style="list-style-type: none"> • Minimum age requirement: 12 years old. • Letter of medical necessity must include documentation and history of other treatments. • Approved for the following documented diagnoses: cervical dystonia, strabismus, or blepharospasm. • Treatment is covered every 3 months, not to exceed 300 units within 90 days. • Not approved for the following uses: primary axillary hyperhidrosis, cosmetic procedures, spasticity. • Prior authorization is required when the medication is obtained through a pharmacy billing through the Point of Sale system. • Initial authorization is given for 6 months. Subsequent authorizations require documentation of patient progress.
Brand Name Medication	<ul style="list-style-type: none"> • Provide details of adverse reaction, allergy, or inadequate response • Authorization is granted for one year. Subsequent authorizations require a telephone call from the physician's office or pharmacy.
Brand Name Schedule II Meds	<ul style="list-style-type: none"> • Documentation from progress notes detailing the patient's allergic skin reaction or adverse reaction. • Authorization is granted for one year. Subsequent authorizations require an updated letter of medical necessity. <p>NOTE: Prior authorizations for brand name medications in this drug class require physician evaluated, charted documentation of an allergic reaction or adverse reaction. Patient complaints of lack of efficacy, such as "patient said", "patient reports", "doesn't work", or "causes nausea" are not acceptable reasons for failure.</p>

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Cancidas	<ul style="list-style-type: none"> Criteria for Invasive Aspergillosis Infection: <ul style="list-style-type: none"> Minimum age: 18 years old. Failure on Amphotericin B OR Documented lab culture showing that aspergillosis is not sensitive to Amphotericin B or itraconazole. Criteria for Candida: <ul style="list-style-type: none"> Minimum age: 18 years old. Diagnosis of esophageal candida, intra-abdominal abscess, peritonitis, or pleural space infections. <ul style="list-style-type: none"> Cultures identifying candida. Also approved for prophylaxis for severely immuno-compromised bone marrow transplant patients with severe graft vs. host disease. Authorizations are granted for 3-month periods and require documentation of lab cultures and continuing symptoms. Children with cancer or bone marrow transplants and adults with bone marrow transplants may receive it as needed following hospitalization.
Celebrex	<ul style="list-style-type: none"> Provide documentation of one of the following diagnoses: <ul style="list-style-type: none"> GERD Barrett's Syndrome Peptic Ulcer Gastro-hypersecretory condition or gastric bleeding caused by other NSAIDS (Documentation from progress notes is required). History of Ulcers Concomitant anticoagulant therapy Failure on 3 other NSAIDS (Documentation from progress notes is required) Prior authorization is not required for age 65+ - see page 3. Analgesia for 10 days will be granted with a telephone call from the physician's office or pharmacy. Initial authorization is for one year - renewals require a telephone request from the physician's office or pharmacy.
Combunox	<ul style="list-style-type: none"> Components must be unavailable separately. A telephone call from the pharmacy or provider to the PA team is required. Authorization is granted for a maximum of 4/day for a 7 day supply. Re-authorization requires a new PA request.
Cytogam	<ul style="list-style-type: none"> Covered for the prophylaxis of cytomegalovirus Physician must provide documentation of transplantation of kidney, lung, liver, pancreas, or heart. Initial authorization is for 6 months - renewals require a telephone request from the physician's office or pharmacy.

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<p>Diabetes Medications</p> <ul style="list-style-type: none"> Byetta Exubera Insulin Pens & Cartridges 	<p>Byetta:</p> <ul style="list-style-type: none"> Minimum age requirement - 17 years old. The patient cannot be using insulin. Byetta cannot be a replacement for insulin. Byetta will only be approved as an adjunct therapy in the treatment of Type II Diabetes. Patient must be taking metformin, a sulfonourea (identify by name) or both OR a TZD (glitazone) alone or in combination with metformin. The patient cannot be in end-stage renal disease or on dialysis. The patient may not have a diagnosis of gastroparesis. Provide information showing a lack of glycemic control. Initial authorization is for 1 year - renewal requires documentation that the patient is stable on Byetta and is not on insulin. <p>Exubera:</p> <ul style="list-style-type: none"> Minimum age requirement - 18 years old. Diagnosis of Type I or Type II Diabetes. Not approved for smokers. Detailed description of medical necessity, including a description of the patient's underlying pulmonary condition. Documentation of why the patient is unable to use short-acting insulin. Approval will not be granted for patient convenience. Is not being used in combination with a short-acting insulin (long acting is OK). Initial authorization is for 1 year - renewals require a letter from the physician showing that the above criteria are still being met. <p>Insulin Pens & Cartridges:</p> <ul style="list-style-type: none"> Medicaid will only pay for the insulin cartridge or pen for those that are legally blind. Initial authorization is for 1 year - renewals require a telephone request from the physician's office or pharmacy.
<ul style="list-style-type: none"> Symlin 	<p>Symlin:</p> <ul style="list-style-type: none"> Is being used for Type I or Type II diabetes as adjunct therapy for patients who use mealtime insulin. Patient has failed desired glucose control despite optimal insulin therapy. Patient is insulin compliant and does regular insulin monitoring. Patient has not had a hypoglycemic incident requiring assistance in the last 6 months. Patient does not have gastroparesis or hypoglycemia. Has HbA less than 9% Initial authorization is 1 year - renewals require a telephone call from the physician's office or pharmacy.
Emsam	<ul style="list-style-type: none"> Physician documentation from charted progress notes of failure with a minimum of three other antidepressants, which may include MAOI. Previous intolerance to a trial of oral MAOI. No concurrent antidepressant therapy. Initial authorization is 1 year - renewals require a telephone call from the physician's office or pharmacy. Non-covered under Non-Traditional Medicaid.

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Enzymes <ul style="list-style-type: none"> Adagen Aldurazyme Aralast Cerezyme Fabrazyme Prolastin / Zemaira 	<p>Adagen:</p> <ul style="list-style-type: none"> Documented diagnosis of Adenosine Deaminase Deficiency. Copy of prescription from physician. Dose must be delivered in pre-filled syringe for exact dosing. Medicaid must be notified of changes in dosage with a copy of new prescription. Authorization is for 1 year - renewals require a telephone call from the physician's office of pharmacy. <p>Aldurazyme:</p> <ul style="list-style-type: none"> Documented and confirmed diagnosis of Hurler and Hurler-Scheie. Confirmed diagnosis is defined as Hurler and Hurler Scheie of mucopolysaccharidosis I (MPS I) in patients with Scheie form who have severe symptoms. Initial authorization is for 6 months - renewals require a telephone call from the physician's office or pharmacy. <p>Aralast:</p> <ul style="list-style-type: none"> Diagnosis of emphysema. History of treatment, including current treatment and past treatment failures. Explanation of condition that demands augmentation with Aralast. Initial authorization is for 6 months - renewals are granted for 1 year periods with documentation of sustained improvement. <p>Cerezyme:</p> <ul style="list-style-type: none"> Documented diagnosis of Gaucher's Disease. Copy of the prescription from physician. Medicaid must be notified of changes in dosage with a copy of new prescription. Initial authorization is for 6 months - renewals are granted in 1 year increments with documentation of significant improvement. <p>Fabrazyme:</p> <ul style="list-style-type: none"> Documented deficient plasma or leukocyte a-galactosidase A (a-gal) OR Documented a-gal deficiency and/or mutation in the a-gal A gene in heterozygous females. Covered only for patients with documented ADA deficiency. Initial authorization is for 6 months - renewals require a telephone request from the physician's office or pharmacy. <p>Prolastin/Zemaira:</p> <ul style="list-style-type: none"> Documented Alpha-1 Antitrypsin deficiency AND Documented Panacinar Emphysema. Must have stopped smoking for at least 30 days, as documented by physician. Initial authorization is for 6 months - renewals require a telephone request from the physician's office or pharmacy.

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<p>Erythropoietins</p> <ul style="list-style-type: none"> Aranesp Procrit Neupogen 	<p>Aranesp:</p> <ul style="list-style-type: none"> Diagnosis of anemia associated with renal failure or chemotherapy. Patient is not on dialysis. Patient does not have a GI bleed. Hematocrit <33% supported by lab work done in the last 3 months (fax copy). Hemoglobin <11% supported by lab work done in the last 3 months (fax copy). Prescribing authority is granted to hematologist, oncologist, nephrologist, and infectious disease specialists, or based upon a consult with one of these specialists. Initial authorization is granted for 6 months - renewals require that the patient not have GI bleeding, not be on dialysis, and lab work in the last 3 months showing hematocrit <39% and Hemoglobin 11-13% (fax copies). <p>Procrit:</p> <ul style="list-style-type: none"> Diagnosis of anemia associated with renal failure, chemotherapy, or HIV. Blood transfusions, allogenic and anemic surgery patients (approve 1 time only). Patient is not on dialysis. Patient does not have a GI bleed. Hematocrit <33% supported by lab work done in the last 3 months (fax copy). Hemoglobin <11% supported by lab work done in the last 3 months (fax copy). Prescribing authority is granted to hematologist, oncologist, nephrologist, and infectious disease specialists, or based upon a consult with one of these specialists. Initial authorization is granted for 6 months - renewals require that the patient not have GI bleeding, not be on dialysis, and lab work in the last 3 months showing hematocrit <39% and Hemoglobin 11-13% (fax copies). <p>Neupogen:</p> <ul style="list-style-type: none"> Documented myelosuppressive chemotherapy, bone marrow transplant, peripheral blood progenitor cell collection, severe chronic neutropenia. Not covered for AIDS, hairy cell leukemia, myelodysplasia, drug-induced congenital agranulocytosis, alloimmune neonatalneutropenia, Hepatitis C. Initial authorization is granted for 6 months - renewals require a telephone request from the physician's office of pharmacy.
<p>Growth Hormone for Adults (AIDS Wasting Syndrome Only)</p>	<ul style="list-style-type: none"> Minimum age - 19 years old. Adult onset AIDS - <u>AIDS Wasting indication only.</u> Body Mass Index is less than 20. Patient must be taking antiretroviral medications. Provide initial height and weight. Rule out other causes of weight loss including hypogonadism (provide testosterone levels for men), opportunistic infections, diarrhea, inadequate nutritional intake, malabsorption, and thyroid abnormalities. Patients must be able to maintain 100% of daily nutritional intake. For patients receiving enteral or parenteral nutrition, the patient must be weight stable for 2 months. Patient must not have an untreated or suspected systemic infection or persistent fever > 101 F during the 30 days prior to evaluation of weight loss. Patient must not have any signs or symptoms of gastrointestinal malabsorption or blockage unless on total parenteral nutrition. Patient must not have active malignancy, except for Kaposi's Sarcoma. Initial authorization is granted for a 60-day trial - renewals require a copy of the current history and physical showing weight gain. With appropriate progress, the patient may receive an additional four weeks of therapy. If the patient continues to show progress, additional prior authorizations are granted in 6 week periods to a maximum of 12 weeks in any 6 months.

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<p>Growth Hormone</p> <ul style="list-style-type: none"> Children 	<ol style="list-style-type: none"> Ages 0 - 18; must start before age 16 Height Stature <5th % on NCHS Growth Chart Growth rate documented 6 months immediately prior Must have: <ul style="list-style-type: none"> -Growth Failure due to: <ul style="list-style-type: none"> -Low Endogenous GH secretion <10ng/ml after provocative stimulation, AND low IGF-I levels as per the testing labs' ranges -documented chronic renal insufficiency up to time of renal transplant, OR -Idiopathic short stature defined by SDS < 2.25 (Humatrope), OR -Turner Syndrome in patients with open epiphysis, OR -Short Bowel Syndrome in patients receiving specialized nutritional support, OR -Panhypopituitarism, OR -Small for gestational age (2 years max coverage): <ul style="list-style-type: none"> -requested before age 3 -normal GH levels or documented GH resistance -catch-up growth not shown before age 2 Completed sleep study for clients with Prader Willi Prescribed by an endocrinologist or with endo consultation
<p>Hepatitis Medications</p> <ul style="list-style-type: none"> Hepsera Rebetron 	<p>Hepsera:</p> <ul style="list-style-type: none"> Diagnosis of hepatitis B. Failure on Epivir. 10mg/day is the maximum approved dose. Initial authorization is granted for 12 weeks - renewals are granted in 12 month cycles with a telephone request from the physician's office or pharmacy. <p>Rebetron:</p> <ul style="list-style-type: none"> Documented diagnosis of hepatitis C. Patient must be severely ill. Initial authorization is granted for 6 months - renewal requests require a letter of medical necessity documenting the current condition.
<p>Hydroxyprogesterone Caproate</p>	<ul style="list-style-type: none"> Approved for the prevention of preterm labor for patients with prior history of preterm delivery. Must be prescribed by an OBGYN. Therapy initiated between 16-23 weeks gestation. Pharmacy provider must submit evidence of compliance with USHP 797 standards for sterile preparation of the injection. Authorization is granted for the duration of the pregnancy. A new PA application must be submitted for each pregnancy.
<p>Increlex</p>	<ul style="list-style-type: none"> Patient age is between 2 and 18. Documented diagnosis of Primary IGF-1 Deficiency. GH and IGF-1 levels are at or below defined limits. Patient does not have cancer. Patient is not on chronic steroid therapy. Patient does not have any uncorrected thyroid deficiencies. Initial authorization is granted for 1 year - renewal requests require the same documentation as the initial request.

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Influenza Medications <ul style="list-style-type: none"> Relenza Tamiflu 	<p>Relenza:</p> <ul style="list-style-type: none"> Minimum age requirement: 7 years old. Diagnosis of Influenza A or Influenza B. Covered only for patients at high risk from diagnosed and documented disease states of immunodeficiency. This includes HIV/AIDS or other diseases of the immune system; long-term radiation treatment; long-term treatment with drugs such as steroids, oncology agents, or immunosuppressive agents; or fragility due to extreme age (greater than 65 years). Prior Approval is limited to one box of 20 amps per year. Treatment must be started within 72 hours of diagnosis. <p>Tamiflu:</p> <ul style="list-style-type: none"> Minimum age requirement: 1 years old. Diagnosis of Influenza A or Influenza B. Covered only for patients at high risk from diagnosed and documented disease states of immunodeficiency. This includes HIV/AIDS or other diseases of the immune system; long-term radiation treatment; long-term treatment with drugs such as steroids, oncology agents, or immunosuppressive agents; or fragility due to extreme age (greater than 65 years). Prior approval is limited to 10 capsules per year. <p style="text-align: center;">OR</p> <ul style="list-style-type: none"> Prophylaxis for Influenza A or B for age 13 and older. Documentation that demonstrates that one other household member or residential member currently has documented influenza A or B. Covered only for patients at high risk from diagnosed and documented disease states of severe cardiopulmonary conditions, immuno-compromised patients, fragility due to extreme age (greater than 65 years). Prior approval is limited to a 7-day course with 14 capsules. TREATMENT MUST BE STARTED WITHIN 72 HOURS OF DIAGNOSIS.
Irritable Bowel Medication <ul style="list-style-type: none"> Amitza 	<p>Amitza:</p> <ul style="list-style-type: none"> Minimum age requirement - 18 years old. Diagnosis of Chronic Idiopathic Constipation. Documented failure within the last 12 months using one fiber laxative and two stimulant laxative products. Drug induced constipation must be ruled out. Initial authorization is granted for 6 months - patient may have a second authorization after a trial off Amitza using other laxatives for at least 30 days. <p style="text-align: center;">OR</p> <ul style="list-style-type: none"> Diagnosis of Irritable Bowel Syndrome with Constipation. Documented failure within the last 12 months using one fiber laxative and one osmotic laxative (magnesium salts or polyethylene glycol based laxatives) Other causes of constipation have been ruled out. Initial authorization is granted for 3 months - patient may have a second authorization after a trial off Amitza using other laxatives for at least 30 days.
Lactulose	<ul style="list-style-type: none"> Documented diagnosis of chronic liver failure, hepatic encephalopathy, chronic portal hypertension, or Spina Bifida. Prior authorization is only required for > 6000ml's per month. This drug will not be approved for use as general laxative for over 6000ml's monthly. Initial authorization is granted for 6 months - renewals require a telephone call from the physician's office or pharmacy.
Lamisil	<ul style="list-style-type: none"> Documented diagnosis of onychomycosis. Coverage will be limited to 16 weeks per calendar year.

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LMWH Derivatives <ul style="list-style-type: none"> Arixtra Fragmin Innohep Lovenox Lovenox (Pregnancy): 	<p>Arixtra:</p> <ul style="list-style-type: none"> Pre-operative for 3 days to stop coumadin prior to surgery. Post-operative for 5 days to achieve therapeutic INR on coumadin. Post-operative prevention of DVT in patients with abdominal surgeries and below (i.e. hip, knee, and ankle not including foot and toes) for a maximum of 10 days. Treatment of acute DVT or PE when administered in conjunction with coumadin, when initial therapy is administered in the hospital. Re-authorization is considered on an individual basis and is based on INR. <p>Fragmin:</p> <ul style="list-style-type: none"> Pre-operative for 3 days to stop coumadin prior to surgery. Post-operative for 5 days to achieve therapeutic INR on coumadin. Post-operative prevention of DVT in patients with abdominal surgeries and below (i.e. hip, knee, and ankle not including foot and toes) for a maximum of 10 days. Treatment of acute DVT or PE when administered in conjunction with coumadin, when initial therapy is administered in the hospital. Re-authorization is considered on an individual basis and is based on INR. Ischemic complications in unstable angina and non-Q-wave MI patients on concurrent aspirin therapy for a maximum of 10 days. Re-authorization is considered on an individual basis and is based on INR. <p>Innohep:</p> <ul style="list-style-type: none"> Documented diagnosis of a DVT or PE. Treatment in conjunction with coumadin regulation and treatment for a maximum of 20 units in 10 days. Re-authorization is considered on an individual basis and is based on INR. Requests may also be made by petition to the DUR Board. <p>Lovenox:</p> <ul style="list-style-type: none"> Pre-operative for 3 days to stop coumadin prior to surgery. Post-operative for 5 days to achieve therapeutic INR on coumadin. Post-operative prevention of DVT in patients with abdominal surgeries and below (i.e. hip, knee, and ankle not including foot and toes) for a maximum of 10 days. Treatment of acute DVT or PE when administered in conjunction with coumadin, when initial therapy is administered in the hospital. Re-authorization is considered on an individual basis and is based on INR. Ischemic complications in unstable angina and non-Q-wave MI patients on concurrent aspirin therapy for a maximum of 10 days. Re-authorization is considered on an individual basis and is based on INR. <p>Lovenox During Pregnancy:</p> <ul style="list-style-type: none"> Past history of DVT/PE OR Active DVT/PE OR Known hypercoagulability.
Myobloc	<ul style="list-style-type: none"> Minimum age requirement: 12 years old. Letter of medical necessity must include documentation and history of other treatments. Approved for a documented diagnosis of cervical dystonia. Treatment is covered every 3 months, not to exceed 10,000 units within 90 days. Prior authorization is required when the medication is obtained through a pharmacy billing through the Point of Sale system. Initial authorization is given for 6 months. Subsequent authorizations require documentation of patient progress.
Multiple Sclerosis Medications <i>(Avonex, Copaxone, Rebif)</i>	<ul style="list-style-type: none"> Documented diagnosis of Multiple Sclerosis. Initial authorization is granted for 1 year. Renewals require a telephone call from the physician's office or pharmacy.

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Oxandrin	<ul style="list-style-type: none"> First 60 day trial period: <ul style="list-style-type: none"> Minimum age requirement - age 19. Adult onset AIDS wasting indication only. BMI is less than 20 - provide current height, weight and BMI. Patient must be taking antiretroviral, documented. Patient must be maintaining a nutritional intake. Authorization after 60 day trial (may approve for an additional 4 months): <ul style="list-style-type: none"> All criteria above remains effective. Weight needs to have been maintained or has increased. If weight has not maintained, it is no longer a benefit. Patient may need to advance to growth hormone. Subsequent authorizations are granted in 6 month periods, and require documentation that the patient's weight has maintained or increased. Provide previous weight and current height.
Proton Pump Inhibitors	<ul style="list-style-type: none"> Once daily dosing does not require an authorization - see page 7. Prilosec OTC does not require a prior authorization for BID dosing. Prior authorizations will be allowed for presenting acute states of GERD, ulcers, or hypersecretory conditions. Documentation required includes a copy of an endoscopy report done within the last two years showing GERD or ulcers, or a copy of a hypersecretory study showing the hypersecretory condition. Initial authorization is granted for two months. BID dosing for longer than 2 months requires special approval from the DUR board.
Provigil	<ul style="list-style-type: none"> Minimum age requirement - age 9. Covered for the following diagnoses: <ul style="list-style-type: none"> Narcolepsy - Amphetamines or Methylphenidate must be tried first. Dose is limited to 400mg daily. Treatment to offset sedation related to multiple sclerosis treatment modalities. Dose is limited to 200mg daily. Daytime somnolence due to obstructive sleep apnea - <u>must</u> be on C-pap. Dose is limited to 200mg daily. Initial authorization is granted for 1 year - renewals require a telephone call from the physician's office or pharmacy.

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Pulmonary Anti-hypertensives <ul style="list-style-type: none"> Flolan Letairis Remodulin Revatio Tracleer Ventavis 	<p>Flolan:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of pulmonary hypertension. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy. <p>Letairis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of pulmonary hypertension. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy. <p>Remodulin:</p> <ul style="list-style-type: none"> Minimum age requirement: 16 years old. Documented diagnosis of pulmonary hypertension. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy. <p>Revatio:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of pulmonary hypertension. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy. <p>Tracleer:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of pulmonary hypertension. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy. <p>Ventavis:</p> <ul style="list-style-type: none"> Minimum age requirement: 18 years old. Documented diagnosis of pulmonary hypertension. Only the drug is paid through the Medicaid Pharmacy Program. The Prodose AAD system must be billed as medical equipment using code K0370 and does not require a PA. Patients of an HMO (other than Select Health) must make arrangements to receive the Prodose AAD system through their HMO. Initial authorization is for 1 year. Re-authorization requires a telephone call from the physician's office or pharmacy.
Qualaquin	<ul style="list-style-type: none"> Documented diagnosis of malaria. One 7 day course of a maximum of 42 tablets is approved with each PA.

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Regranex	<ul style="list-style-type: none"> • Rule out venous ulcers and/or arterial ulcers. • Patient must be diabetic, either type I or type II. • Not covered for diabetic ulcers above the ankle. • Patient must have stage III or IV diabetic foot or ankle ulcer as per the International Association of Enterostomal therapy guide to chronic wound staging 1989. • Not a benefit for patients in long term care facilities, unless that patient is admitted from home or hospital with a pre-existing diabetic ulcer of the lower extremity. LTCF must submit a copy of skin assessment report made within 24 hours of admission. • The client must have had a documented failure on a 60 day regimen of good ulcer care that includes but is not limited to: <ol style="list-style-type: none"> 1. Initial complete sharp debridement. 2. A non-weight bearing regimen. 3. Systemic treatment for wound-related infections. 4. Moist saline dressing changes twice daily. 5. Additional debridement If necessary. • The subcutaneous ulcer may not exceed 3cm in diameter or total surface of 9.42cm² (size and shape must be documented). • Total contact casting is an available method of treatment and must be considered and rejected before Regranex is to be considered. • Initial authorization may be granted for 8 weeks and 15-30gm - renewal requires a second PA application demonstrating a 30% reduction in ulcer size. • Treatment is limited to 60gm of Regranex.
Relistor	<ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Diagnosis of opioid-induced constipation. • Patients must be receiving opioid therapy as part of a palliative care regimen for advanced illness. • Rule out mechanical GI obstruction. • Documentation of trial and failure of conventional laxative therapy. • Initial authorization is granted for 4 months. Renewals may be granted with a telephone call from the physician's office or pharmacy.
Renal Cell Carcinoma Meds <ul style="list-style-type: none"> • Nexavar • Sutent 	<p>Nexavar:</p> <ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Diagnosis of advanced renal cell carcinoma. • Initial authorization is granted for 400mg BID until no benefit or side-effects are intolerable - renewal requests are granted with a telephone call from the physician's office or pharmacy. • Nexavar is available only through 5 specialty pharmacies via mail-order: Caremark, Curascript, Accredo, Pharmicare, or McKesson Specialty. <p>Sutent:</p> <ul style="list-style-type: none"> • Minimum age requirement 18 years old. • Diagnosis and documentation of advanced renal cell carcinoma. • History of other treatments, including documented disease progression on or intolerance to Gleevec. • Initial authorization is granted for 50mg daily, 4 weeks on and 2 weeks off. Dose increases or reductions by 12.5mg increments approved as needed or tolerated. Renewals may be granted with a telephone call from the physician's office or pharmacy.

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Drugs Requiring Prior Authorization	
Restasis	<ul style="list-style-type: none"> Approved for the following diagnoses (ICD.9: <ul style="list-style-type: none"> 370.20 (Superficial keratitis, unspecified) 370.21 (Punctate keratitis) 370.33 (Keratoconjunctivitis sicca, not specified as Sjogren's disease) 710.2 (Sicca syndrome - Sjogren's disease) Documentation requirements for the above diagnoses: <ol style="list-style-type: none"> Diagnosis. Documented fluorescein test. Request from ophthalmologist or with documented ophthalmologist consult. Prior approval for the above diagnoses is granted for 1 year - renewals require a new PA request. <p style="text-align: center;">OR</p> <ul style="list-style-type: none"> Documented corneal transplant (ICD.9 V 42.5) Initial authorization is granted for 1 year - renewals granted with a telephone call from physician's office or pharmacy.
Retinoids <ul style="list-style-type: none"> Panretin 	Panretin: <ul style="list-style-type: none"> Initial 30-day trial period: <ul style="list-style-type: none"> Diagnosis of cutaneous lesions caused by Kaposi's Sarcoma Documentation of primary number of KS lesions, estimated total square centimeters, number of lesions flat on baseline, and number of lesions raised on baseline. Systemic anti-KS therapy is not yet required. Retin-A 0.1% gel has been tried for a period of 60 or more days, and there was less than a 25% improvement of both partial response area and partial response height. 60 day treatment period: <ul style="list-style-type: none"> Patient must sustain partial response defined as a 50% or more improvement from base line. Documentation of primary number of KS lesions, estimated total square centimeters, partial response area, and partial response height. Re-authorization may be granted for additional treatment of 60 day periods with continued improvement documented as above.
<ul style="list-style-type: none"> Retin-A 	Retin-A: <ul style="list-style-type: none"> Diagnosis of cutaneous lesions caused by Kaposi's Sarcoma. Pre-pancretin use. Documentation of primary number of KS lesions, estimated total square centimeters, number of lesions flat on baseline, and number of lesions raised on baseline. Systemic anti-KS therapy is not yet required. Initial authorization is granted for a 60-day trial period. Re-authorization is given for 6 month periods with documentation indicating that the patient has had at least a 25% improvement from baseline.
Risperdal Consta	<ul style="list-style-type: none"> Minimum age requirement - 18 years old. Documentation of patient diagnosis (open for the same ICD-9 codes as oral risperdal). Documentation that the patient is unresponsive to conventional treatment. Documentation that the patient is non-compliant with previous treatment modalities. Drug must be administered in a clinic or physician office, NOT approved for nursing homes or group homes. The initial prior approval must be obtained by a prescriber associated with a capitated mental health plan. Approved only for one injection at two-week intervals. Initial authorization is granted for one year - renewal requests require a telephone call from the physician's office.

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Drugs Requiring Prior Authorization	
Selzentry	<ul style="list-style-type: none"> • Minimum age requirement - 16 years old. • Documentation of co-receptor tropism assay test indicating CCR5-tropic HIV-1 infection. • Documentation of optimized background therapy for the treatment of HIV-1 infection. • Initial authorization is for 12 months - renewal requests require a telephone call from the physician's office or pharmacy.
Soliris	<ul style="list-style-type: none"> • Documented diagnosis of paroxysmal nocturnal hemoglobinuria. • Documented failure of or intolerance to other PNH treatments, including transfusion. • Review by the DUR Board.
Somavert	<ul style="list-style-type: none"> • Documented acromegaly. • Documentation showing inadequate response to either transsphenoidal adenomectomy or radiotherapy or both. • Documented trial on at least one dopamine agonist such as cabergoline or bromocriptine. • Documentation that the patient has been evaluated for a somatostatin analogue such as octreotide acetate. • Initial authorization is granted for one year - renewal requests require a telephone call from the physician's office of pharmacy.
Stimulants <ul style="list-style-type: none"> • Adult ADHD Stimulants 	Adult ADHD Stimulants: <ul style="list-style-type: none"> • Documented Diagnosis of one of the following: ADD, ADHD, narcolepsy, organic brain syndrome, traumatic brain injury, treatment resistant depression, mental retardation (if the patient exhibits injurious behavior, is hyperactive, or both), severe sedation due to psychotropic or chemotherapeutic medications. • Letter of medical necessity stating current treatment and situation. • Depression diagnosis requires a description of treatment history and failures. • Adult ADD/ADHD diagnosis requires a copy of the testing that has been done to make the diagnosis of adult ADD/ADHD. Acceptable testing for adult ADD/ADHD includes a psychiatric evaluation, Wender Utah Rating Scale scoring 46 or greater, documentation of the criteria that have been met from the DSM IV manual. • Statement documenting and substance abuse problems past or present, or a statement indicating no substance abuse history. • Initial authorization may be granted for one year - renewal requests require an updated medical necessity <u>and</u> an updated substance abuse statement.
Synagis	<ul style="list-style-type: none"> • Infants of 28 week gestation may receive Synagis prophylactically during the first year of life. • Infants of 29-35 weeks gestation may receive Synagis prophylactically during the 1st to 6th month of life. • Any children under 24 months may receive Synagis if they have either <ol style="list-style-type: none"> 1. Clinical diagnosis of Broncho Pulmonary Dysplasia (BPD) requiring ongoing medical treatment OR 2. Hemodynamically significant Congenital Heart Disease (CHD) requiring ongoing treatment. • Criteria for coverage through a pharmacy: <ol style="list-style-type: none"> 1. Be home bound; and, 2. The pharmacy must bill using correct NDC numbers. • Synagis is not available to any child with active RSV. • The Utah Medicaid Synagis season is for a 6 month period beginning November 1. • A total of 5 immunizations during this 6 month period will be approved, except when the patient begins the immunizations late in the season. • A child who has started the series and then turns 2 may continue to a total of 5 immunizations or to the end of the season, whichever comes first. • No approval will be given to a child 24 months of age or older. • Physicians who provide the vaccine in the office should use code 90378 and the appropriate administration code for reimbursement.

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Drugs Requiring Prior Authorization	
Tasigna	<ul style="list-style-type: none"> • Minimum age requirement: 18 years old. • Diagnosis of chronic myelogenous leukemia. • Documented intolerance or resistance to therapy that includes Gleevec. • Initial authorization is for 1 year. • Re-authorization requires a telephone call from the physician's office or pharmacy.
Trizivir	<ul style="list-style-type: none"> • Documented failure of all three medications (Abacavir, Lamivudine, and Zidovudine) individually. • Initial authorization may be granted for 1 year - renewal requests require a telephone call from the physician's office or pharmacy.
Tykerb	<ul style="list-style-type: none"> • Minimum age: 18 years old. • Diagnosis of advanced or metastatic breast cancer whose tumor overexpresses HER2. • Prior therapy, including an anthracycline, a taxane, and trastuzumab. • To be given in combination with capecitabine. • Prior authorization is given for 1 year - renewal requires an updated letter of medical necessity.
Vectibix	<ul style="list-style-type: none"> • Minimum age - 18 years old. • Diagnosis of metastatic colorectal cancer. • Disease progression on or following fluoropyrimidine-, oxplatin-, and irinotecan-containing chemotherapy regimens. • Initial authorization may be granted for 1 year - renewal requires an updated letter of medical necessity.
Vivitrol	<ul style="list-style-type: none"> • Diagnosis of alcohol abuse. • Negative urine screen for opioids or passed naloxone challenge. • Description of the psychosocial support to be received by the patient, as indicated by chart notes or a brief letter of medical necessity. • Negative screen for liver problems.
Xanax XR	<ul style="list-style-type: none"> • Documentation of failure on a 6-8 week trial of short-acting oral alprazolam within the last 6 months. • Initial authorization may be granted for 1 year - renewal requests require a telephone call from the physician's office or pharmacy. • Non-covered under Non-Traditional Medicaid.
Xibrom	<ul style="list-style-type: none"> • Prior trial of any indicated medication. • Approved for one bottle for a 2 week period following procedure or surgery.
Xolair	<ul style="list-style-type: none"> • Minimum age requirement - 12 years of age. • Diagnosis of moderate to severe persistent asthma of at least 1 year duration OR • Diagnosis of allergic conjunctivitis and rhinitis, atopic dermatitis, and food allergy. • Documented positive skin test reaction to a perennial aeroallergen. • Documentation showing symptoms are inadequately controlled with inhaled corticosteroids and beta-agonists. • Documented forced expiratory volume in one second (FEV1) < 70% predicted (It is possible for severe asthmatics to be better than 70% FEV1 on rescue therapy, but they would still struggle with optimum maintenance therapy. In such severe cases, Xolair would be considered for approval). • Documented pulmonologist or allergist consultation within the last 60 days. • Body weight > 30kg and < 150kg. • Baseline total serum IgE >30 and < 700 IU/ml. • Patient prescription claim history must demonstrate routine use of inhaled corticosteroids for a 90 day period. • Initial authorization may be granted for 6 months - renewal requests require a telephone call from the physician's office or pharmacy.

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Drugs Requiring Prior Authorization	
Xolegel	<ul style="list-style-type: none"> • Minimum age: 12 years old. • Documented trial and failure of a generic formulation of topical ketoconazole within the last 12 months. • Prior authorization is given for 6 months - renewal requests require a telephone call from the physician's office or pharmacy.
Xyrem	<ul style="list-style-type: none"> • Age requirement - 18 to 65 years old. • Documented cataplexy associated with narcolepsy. • Documentation ruling out concomitant use of sedative-hypnotics. • Maximum dose is 9gm/day • Initial authorization may be granted for 1 year - renewal requests require a telephone call from the physician's office or pharmacy.
Zavesca	<ul style="list-style-type: none"> • Minimum age requirement - 18 years old. • Diagnosis of moderate type I Gaucher's disease. • Documentation that enzyme replacement therapy has failed. • Platelet count > 50k/ul (FAX a copy of the lab work) • Written consultation with a trained specialist (hematologist or geneticist) • Cumulative limit of 90 capsules in 30 days. • Initial authorization period may be granted for 1 year - renewal requests require a telephone call from physician's office or pharmacy
Ziana	<ul style="list-style-type: none"> • Age requirement - 12-19 years old. • Patient must try and fail on a combination of both generic tretinoin gel and clindamycin gel. • Initial authorization may be granted for 1 year - renewal requires an updated letter of medical necessity.

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Utah Medicaid Pharmacy Services

Request for Prior Authorization

Request Date_____

Patient Name_____

Patient DOB ____/____/____ Medicaid ID #_____

Social Security Number (if Medicaid ID# unknown)_____

Pharmacy Name_____

Pharmacy Telephone Number_____

Drug Name and Strength_____

Dosage_____

Prescriber Name_____ National Provider ID_____

Prescriber Telephone#_____ Prescriber Fax #_____

Diagnosis:

Date of Diagnosis:

Attach Supporting Documentation

Total Pages_____

Instructions for submitting this PA Request

- Prior Authorizations are only accepted by fax.
- The form may be completed electronically or *legibly* by hand.
- It is not mandatory to use this cover letter; however all of the information requested on this form is necessary before a prior authorization request can be initiated.
- Fax all necessary documentation to the Medicaid Prior Authorizations Team at the following numbers: (801)536-0964 or (801)536-0960 or (801)536-0959.

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